Nov 6, 2018

TREND Statement Checklist

| Paper | Item | n Descriptor | | Reported? | |
|----------------------|------|--|------|-----------|--|
| Section/ Topic | No | | | Pg # | |
| Title and Abst | ract | | | | |
| Title and | 1 | Information on how unit were allocated to interventions | 1/ | 2 | |
| Abstract | | Structured abstract recommended | V | 2 | |
| | | Information on target population or study sample | V | 2 | |
| Introduction | | | | | |
| Background | 2 | Scientific background and explanation of rationals | | 111 | |
| background | | Scientific Suckground and explanation of rationale | | 4 | |
| | | Theories used in designing behavioral interventions | V | 8-0 | |
| Methods | | | | | |
| Participants | 3 | Eligibility criteria for participants, including criteria at different levels in | V | 1 | |
| | | recruitment/sampling plan (e.g., cities, clinics, subjects) | V | (1 | |
| | | Method of recruitment (e.g., referral, self-selection), including the | V | | |
| | | sampling method if a systematic sampling plan was implemented | V | 15 | |
| | | Recruitment setting | V | 15 | |
| | | Settings and locations where the data were collected | | 15 | |
| Interventions | 4 | Details of the interventions intended for each study condition and how | | | |
| | | and when they were actually administered, specifically including: | | | |
| | | O Content: what was given? | V | 12-1 | |
| | | Delivery method: how was the content given? | | (2-1 | |
| | | Unit of delivery: how were the subjects grouped during delivery | | (2-1 | |
| | | Deliverer: who delivered the intervention? | | 12-1 | |
| | | Setting: where was the intervention delivered? | 1 | 12-1 | |
| | | Exposure quantity and duration: how many sessions or episodes | or | 10 | |
| | | events were intended to be delivered? How long were they | 1/ | 10 | |
| | | intended to last? | | 12-10 | |
| | | o Time span: how long was it intended to take to deliver the | ./ | 10 1 | |
| | | intervention to each unit? | ~ | 101 | |
| | | Activities to increase compliance or adherence (e.g., incentives) | V | (2-14 | |
| Objectives | 5 | Specific objectives and hypotheses | V | 9-11 | |
| Outcomes | 6 | Clearly defined primary and secondary outcome measures | (2) | 1-11 | |
| | | Methods used to collect data and any methods used to enhance the | | | |
| | | quality of measurements | 17 | 15-1 | |
| | | Information on validated instruments such as psychometric and biometr | ic , | | |
| | | properties | MA | | |
| Sample Size | 7 | How sample size was determined and, when applicable, explanation of a | ny | | |
| | | interim analyses and stopping rules | WA | 11-11 | |
| Assignment Method | 8 | Unit of assignment (the unit being assigned to study condition, e.g., | | | |
| | | individual, group, community) | V | 1 | |
| | | Method used to assign units to study conditions, including details of any | | | |
| | | restriction (e.g., blocking, stratification, minimization) | C | (5 | |
| | | Inclusion of aspects employed to help minimize potential bias induced do | 10 | | |
| | | to non-randomization (e.g., matching) | | (5 | |

TREND Statement Checklist

| | | CHECKIIST | | | |
|-------------------------|-----|---------------------|--|-----|-------|
| Blinding (masking) | 9 | those a | er or not participants, those administering the interventions, and assessing the outcomes were blinded to study condition assignment; atement regarding how the blinding was accomplished and how it sessed. | UlA | |
| Unit of Analysis | 10 | interve | ntion of the smallest unit that is being analyzed to assess ention effects (e.g., individual, group, or community) | ~ | 15 |
| | | metho | nit of analysis differs from the unit of assignment, the analytical d used to account for this (e.g., adjusting the standard error tes by the design effect or using multilevel analysis) | N/A | |
| Statistical Methods | 11 | Statisti outcon | cal methods used to compare study groups for primary methods ne(s), including complex methods of correlated data | V | 10 |
| | | analyse | cal methods used for additional analyses, such as a subgroup es and adjusted analysis | MA | |
| | | | ds for imputing missing data, if used | NIA | ļ |
| | | • Statisti | cal software or programs used | V | 17 |
| Results | | | | | |
| Participant flow | 12 | • Flow or | participants through each stage of the study: enrollment, | T | |
| , | | | nent, allocation, and intervention exposure, follow-up, analysis (a | ~ | fig |
| | | diagran | n is strongly recommended) | | 1 |
| | | 0 | Enro Iment: the numbers of participants screened for eligibility, | + | |
| | | | found to be eligible or not eligible, declined to be enrolled, and enrolled in the study | ~ | fig (|
| | | 0 | Assignment: the numbers of participants assigned to a study condition | V | fig 1 |
| | | 0 | Allocation and intervention exposure: the number of participants assigned to each study condition and the number of participants who received each intervention | ~ | fin (|
| | | 0 | Follow-up: the number of participants who completed the follow-up or did not complete the follow-up (i.e., lost to follow-up), by study condition | / | fig 1 |
| | | 0 | Analysis: the number of participants included in or excluded from the main analysis, by study condition | ~ | fig 1 |
| | | reasons | | ~ | 17 |
| Recruitment | 13 | | efining the periods of recruitment and follow-up | V | 15 |
| Baseline Data | 14 | study c | e demographic and clinical characteristics of participants in each ondition | MA | |
| | | disease | e characteristics for each study condition relevant to specific prevention research | NIA | |
| | | and by | comparisons of those lost to follow-up and those retained, overall study condition | NIA | |
| | 4.5 | of inter | | N/A | |
| Baseline equivalence | 15 | Data on to contr | study group equivalence at baseline and statistical methods used ol for baseline differences | N/A | |

TREND Statement Checklist

| 16 | T- N. I. S. VI. | | _ |
|----|--|---|---|
| 10 | study condition, particularly when the denominators change for different outcomes; statement of the results in absolute numbers when feasible | ~ | 20 |
| | not, description of how non-compliers were treated in the analyses | NIA | |
| 17 | For each primary and secondary outcome, a summary of results for each estimation study condition, and the estimated effect size and a confidence interval to indicate the precision | V | 19 tables |
| | Inclusion of null and negative findings | V | 20 |
| | Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any | V | 10 |
| 18 | Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory | NA | |
| 19 | Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals) | N/A | |
| | | | |
| 20 | Interpretation of the results, taking into account study hypotheses, sources of potential bias, imprecision of measures, multiplicative analyses, and other limitations or weaknesses of the study | ~ | 11 |
| | Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations | V | 21 |
| | fidelity of implementation | V | 22 |
| | | ~ | 22 |
| 21 | Generalizability (external validity) of the trial findings, taking into account the study population, the characteristics of the intervention, length of follow-up, incentives, compliance rates, specific sites/settings involved in the study, and other contextual issues | V | 21-2 |
| 22 | General interpretation of the results in the context of current evidence and current theory | / | 22 |
| | 18 19 20 | study condition, particularly when the denominators change for different outcomes; statement of the results in absolute numbers when feasible Indication of whether the analysis strategy was "intention to treat" or, if not, description of how non-compliers were treated in the analyses For each primary and secondary outcome, a summary of results for each estimation study condition, and the estimated effect size and a confidence interval to indicate the precision Inclusion of null and negative findings Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals) Interpretation of the results, taking into account study hypotheses, sources of potential bias, imprecision of measures, multiplicative analyses, and other limitations or weaknesses of the study Discussion of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations Discussion of the success of and barriers to implementing the intervention, fidelity of implementation Discussion of research, programmatic, or policy implications Generalizability (external validity) of the trial findings, taking into account the study population, the characteristics of the intervention, length of follow-up, incentives, compliance rates, specific sites/settings involved in the study, and other contextual issues General interpretation of the results in the context of current evidence | study condition, particularly when the denominators change for different outcomes; statement of the results in absolute numbers when feasible Indication of whether the analysis strategy was "intention to treat" or, if not, description of how non-compliers were treated in the analyses For each primary and secondary outcome, a summary of results for each estimation study condition, and the estimated effect size and a confidence interval to indicate the precision Inclusion of null and negative findings Inclusion of results from testing pre-specified causal pathways through which the intervention was intended to operate, if any Summary of other analyses performed, including subgroup or restricted analyses, indicating which are pre-specified or exploratory Summary of all important adverse events or unintended effects in each study condition (including summary measures, effect size estimates, and confidence intervals) Interpretation of the results, taking into account study hypotheses, sources of potential bias, imprecision of measures, multiplicative analyses, and other limitations or weaknesses of the study Interpretation of results taking into account the mechanism by which the intervention was intended to work (causal pathways) or alternative mechanisms or explanations Discussion of research, programmatic, or policy implications Discussion of research, programmatic, or policy implications Generalizability (external validity) of the trial findings, taking into account the study population, the characteristics of the intervention, length of follow-up, incentives, compliance rates, specific sites/settings involved in the study, and other contextual issues General interpretation of the results in the context of current evidence |

From: Des Jarlais, D. C., Lyles, C., Crepaz, N., & the Trend Group (2004). Improving the reporting quality of nonrandomized evaluations of behavioral and public health interventions: The TREND statement. American Journal of Public Health, 94, 361-366. For more information, visit: http://www.cdc.gov/trendstatement/